



DEPARTMENT OF HEALTH AND HUMAN SERVICES

Purged DJW  
Food and Drug Administration  
Atlanta District Office m3287n  
HFI-35  
60 Eighth Street, N.E.  
Atlanta, Georgia 30309

December 14, 1999

VIA FEDERAL EXPRESS

Beth Golson  
Chief Technologist  
Metrolina Women's Breast Center  
Park Place Professional Center  
441 N. Wendover Road  
Charlotte, NC 28211

Inspection ID: 1250700005

WARNING LETTER

Dear Ms. Golson:

Your facility was inspected on 11/30/99 by a representative of the North Carolina Department of Environment & Natural Resources (DENR), Division of Radiation Protection, acting on behalf of the Food and Drug Administration (FDA). This inspection revealed that your facility failed to comply with certain Quality Standards for Mammography as specified in Title 21, Code of Federal Regulations (CFR), Part 900.12, as follows:

The system to communicate results is not adequate for your site because there is no system in place to provide timely medical reports.

The specific deficiency noted above appeared under the Level 1 heading on your MQSA facility Inspection Report, which was issued at the close of the inspection. This deficiency may be symptomatic of serious underlying problems that could compromise the quality of mammography at your facility.

In addition, your response should address the Level 2 noncompliance items that were listed on the inspection report provided to you at the close of the inspection. The Level 2 noncompliance items are:

There is no written procedure for handling consumer complaints at your site.

Your firm has no written procedure for infection control.

1 of 6 random reports reviewed did not contain an assessment category.

It is your responsibility to ensure adherence to each requirement of the Mammography Quality Standards Act of 1992 (MQSA) and FDA's regulations. You are responsible for investigating and determining the causes of the deficiencies that the inspection identifies and promptly initiating permanent corrective actions.

If you fail to promptly correct these deficiencies, FDA may, without further notice, initiate regulatory action. Under MQSA, FDA may:

- impose civil money penalties on a facility of up to \$10,000 for each failure to substantially comply with, or each day of failure to substantially comply with, the Standards.
- suspend or revoke a facility's FDA certificate for failure to comply with the Standards.
- seek an injunction in federal court to prohibit any mammography activity that constitutes a serious risk to human health.

Please note that FDA regulations do not preclude a State from enforcing its own State mammography laws and regulations. In some cases, these requirements may be more stringent than FDA's. When you plan your corrective action(s), therefore, you should consider the more stringent State requirements, if any.

Within 15 working days after receiving this letter, you should notify FDA in writing of:

- the specific steps you have taken to correct all of the violations noted in this letter;
- each step your facility is taking to prevent the recurrence of similar violations;
- equipment settings (including technique factors), raw test data, and calculated final results, where appropriate; and
- sample records that demonstrate proper recordkeeping procedures, if the noncompliances that were found relate to quality control or other records. (Note: Patient names or identification should be deleted from any copies submitted.)

If your facility is unable to complete the corrective action within 15 working days, you should state the reason for the delay and the time within which corrections will be completed.

Please send the original copy of your response to:

U.S. Food and Drug Administration  
Compliance Branch  
60 8<sup>th</sup> St., NE  
Atlanta, GA 30309

With a copy to:

North Carolina DENR  
Division of Radiation Protection  
3825 Barrett Drive  
Raleigh, NC 27609-7221

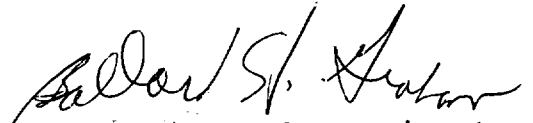
and

Thomas Clarida  
U.S. Food and Drug Administration  
5701 Executive Center Drive, Suite 104  
Charlotte, NC 28212

(NOTE: If phantom image is required for corrective action, please submit original to North Carolina DENR, Division of Radiation Protection.)

You may choose to address both FDA and state requirements in your response. If you have any questions regarding this letter or how to ensure you are meeting MQSA standards, please call Thomas Clarida at 704-344-6116.

Sincerely yours,



Ballard H. Graham, Director  
Atlanta District